9

outlet port 49 of the third test strip dock 48 until the third test strip 80 is received within the third test strip dock 48. Furthermore, the first end 92 of the fourth test strip 90 can be placed within the fourth outlet port 51 of the fourth test strip dock 50 until the fourth test strip 90 is received within the 5 fourth test strip dock 50.

Referring to FIG. 3, upon selecting a site, a clinician can adhere the adhesive on the bottom surface 38 of the lower portion 34 of the housing 30 onto a skin surface S of a patient where a blood sample will be accessed over a selected sampling site as shown in FIG. 3.

Referring to FIGS. 3-5, a user or an operator may then apply pressure to the dome-shaped surface 36 of the upper portion 32 of the housing 30 to actuate the upper portion 32 from the undeformed position (FIGS. 3 and 6) to the 15 deformed position (FIGS. 4 and 5). Actuation of the upper portion 32 from the undeformed position (FIGS. 3 and 6) to the deformed position (FIGS. 4 and 5) moves the lancet 14 from the pre-actuation position (FIGS. 3 and 6) to the puncturing position (FIGS. 4 and 5) thereby causing the lancing of 20 the skin surface S of the patient by the puncturing end 112 of the lancet 14 as shown in FIG. 5. When the upper portion 32 of the housing 30 is depressed, the puncturing end 112 of the lancet 14 cuts into the skin surface S of the patient's body and capillary blood begins to flow into the inlet port 40 of the 25 housing 30.

After lancing and release of the pressure for the lancing action, the domed shape of the upper portion 32 begins to relax and returns to its original shape or undeformed position. This return of the upper portion 32 to its undeformed position 30 creates a gentle vacuum during the process that helps to draw out the capillary blood through the inlet port 40 and to the reservoir or fluid channel 42 of the housing 30. With the upper portion 32 of the housing 30 returned to its undeformed position and with the blood sample 16 received within the 35 reservoir 42 as shown in FIG. 6, the blood sample 16 is directed to flow through the reservoir 42 and to the test strip docks such that the test elements 60, 70, 80, 90 each receive a portion of the blood sample 16. In one embodiment, the reservoir or fluid channel 42 may include microfluidic path- 40 ways integrated therein to direct the blood sample 16 to the test strips 60, 70, 80, 90 by capillary action. The blood receiving cavities 66, 76, 86, 96 are adapted to receive a portion of the blood sample therein.

Referring to FIG. 7, with the blood sample transferred to 45 the test strips or test elements 60, 70, 80, 90, a user or operator may then grasp the second end 64 of the first test strip 60, for example, and pull the first test element 60 from the first test strip dock 44 of the housing 32. Next, referring to FIG. 8, the collected blood sample within the first test strip 60 is trans- 50 ferred to the test strip reader 202 of the point-of-care testing device 200 to analyze the collected blood and obtain test results for tests such as glucose, cholesterol, or other blood sample results. Referring to FIG. 8, the receiving port 204 of the point-of-care testing device 200 allows for the closed 55 transfer of a blood sample from the first test element 60 to the point-of-care testing device 200. In one embodiment, the viewing window 206 of the test strip reader 202 may indicate to an operator desired information. For example, the viewing window 206 may indicate what the point-of-care testing 60 device 200 is analyzing the blood sample for.

In one embodiment, a portion of the blood sampling device 10 contains a sample stabilizer to promote efficient mixing with the blood sample. The sample stabilizer, can be an anti-coagulant, or a substance designed to preserve a specific 65 element within the blood such as, for example, RNA, protein analyte, or other element. In one embodiment, the sample

10

stabilizer is provided within the inlet port 40 of the housing 30 of the blood sampling device 10 and/or any area where a primary blood sample is collected. In another embodiment, the sample stabilizer is provided in a portion of any of the reservoirs or fluid channels 42 of the housing 30 of the blood sampling device 10 and/or along any portion of a liquid path that the blood sample travels. In other embodiments, the sample stabilizer may be provided in any of the test strip docks of the blood sampling device 10. In one embodiment, each of the fluid channels, reservoirs, test strip docks, and/or each of the test elements could each include a different sample stabilizer. In this manner, a single blood sample could be used for a variety of different tests, each of which could introduce an appropriate, and potentially unique, sample stabilizer for a desired use. A blood sampling device of the present disclosure provides flexibility in the nature of the additives and/or sample stabilizers introduced for a blood sample.

Referring to FIG. 5, upon actuation of the lancet 14 to puncture the skin surface S, no or minimal blood will seep between the stick site and the housing 30 of the blood sampling device 10, and, importantly, any seeped blood will not subsequently enter the blood sampling device 10.

Collection using the blood sampling device 10 of the present disclosure allows "closed system" capillary blood collection with the following advantages, such as avoiding exposure of the operator or patient to the blood and avoiding exposure of the blood to atmosphere and potentially better preservation of in vivo blood gas concentrations. Also, the blood sampling device 10 of the present disclosure incorporates the concepts of lancing, blood collection, and multiple test strip collection. The blood sampling device 10 of the present disclosure allows for a blood sample to be collected on a plurality of test elements simultaneously. For example, the blood sampling device 10 allows for only a single stick on a patient and collection of a blood sample into multiple test strips for a point-of-care testing device. This ability to collect a single blood sample into multiple test strips reduces the anxiety and discomfort to the patient.

While this disclosure has been described as having exemplary designs, the present disclosure can be further modified within the spirit and scope of this disclosure. This application is therefore intended to cover any variations, uses, or adaptations of the disclosure using its general principles. Further, this application is intended to cover such departures from the present disclosure as come within known or customary practice in the art to which this disclosure pertains and which fall within the limits of the appended claims.

What is claimed is:

- 1. A biological fluid sampling device, comprising:
- a housing having an inlet port, a reservoir disposed within the housing and in fluid communication with the inlet port, and a first cavity in fluid communication with the reservoir;
- a first test element removably receivable within the first cavity, wherein the first test element is a test strip; and
- a puncturing element, a portion of which is disposed within the housing and adapted for movement between a preactuated position wherein the puncturing element is retained within the housing and a puncturing position wherein the puncturing element extends through the inlet port of the housing.
- 2. The biological fluid sampling device of claim 1, wherein the biological fluid sampling device is adapted to receive a blood sample.